

INFORMATION DISCLOSURE CITATION (Use several sheets if necessary)				Attorney Docket No. 056291-5215		Application No. 10/554,202	
PTO Form 1449 December 4, 2008				Applicants: Robert Hugh BRADBURY et al.		Filing Date: October 24, 2005	
				Group Art Unit: 1624			
U.S. PATENT DOCUMENTS							
Initial	Document No.	Date	Name	Class	Sub-Class	Filing Date	
	1. US 2003/0186995	October 2, 2003	Kath et al.				
	2. US 2004/0048880	March 11, 2004	Himmelsbach et al.				
ALL REFERENCES CONSIDERED EXCEPT WHERE LINED THROUGH. /T.N./							
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	Document No.	Date	Country	Class	Sub-Class	Translation	
	3. CA 2476888	October 9, 2003	Canada				
	4. CA 2543649	May 12, 2005	Canada				
	5. WO 01/21596	March 29, 2001	WIPO				
	6. WO 2004/046101	June 3, 2004	WIPO				
	7. WO 2005/041973	May 12, 2005	WIPO				
	8. WO 2005/097134	October 20, 2005	WIPO				
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OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Pages, etc.)							
9.	Ballard et al. "Developing a small molecule erbB2 inhibitor: challenges with optimising DMPK properties" Poster - Presented at DMDG Cambridge (February 6, 2008) .						
10.	Ballard et al. "Neutral 5-substituted 4-anilinoquinazolines as potent, orally active inhibitors of erbB2 receptor tyrosine kinase" Bioorg Med Chem Lett. 17(22):6326-6329 (2007) .						
11.	Barlaam et al. "A new series of neutral 5-substituted 4-anilinoquinazolines as potent, orally active inhibitors of erbB2 receptor tyrosine kinase" Bioorganic & Medicinal Chemistry Letters 18(2):674-678 (2008) .						
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17.	Gaul et al. "Discovery and Biological Evaluation of Potent Dual ErbB-2/EGFR Tyrosine Kinase Inhibitors: 6-Thiazolylquinazolines" Bioorganic & Medicinal Chemistry Letters 13(4):637-640 (2003) .						
18.	Harris et al. "Systematic variation of a key quinazoline core" Presented at the XXII European Colloquium on Heterocyclic Chemistry (XXII ECHC-2006) Bari, Italy, September 2-6, 2006 .						
19.	Hennequin et al. "N-(5-chloro-1,3-benzodioxol-4-yl)-7-[2-(4-methylpiperazin-1-yl)ethoxy]-5-(tetrahydro-2H-pyran-4-yloxy)quinazolin-4-amine, a novel, highly selective, orally available, dual-specific c-Src/Abl kinase inhibitor" J Med Chem. 49(22):6465-6488 (2006) .						
20.	Jani et al. "Discovery and pharmacologic characterization of CP-724,714, a selective ErbB2 tyrosine kinase inhibitor" Cancer Research 67(20):9887-9893 (2007) .						
Examiner		/Tamthom Truong/		Date Considered		05/07/2009	
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